

ICF International / Laboratory Data Consultants

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MEMORANDUM

TO:

Chris Lichens, Remedial Project Manager

Site Cleanup Section 4, SFD-7-4

THROUGH:

Rose Fong, ESAT Task Order Manager (TOM)

Quality Assurance (QA) Program, MTS-3

FROM:

Doug Lindelof, Data Review Task Manager

Region 9 Environmental Services Assistance Team (ESAT)

ESAT Contract No.: EP-W-06-041

Technical Direction Form No.: 00105074 Amendment 3

DATE:

August 23, 2007

SUBJECT:

Review of Analytical Data, Tier 2

Attached are comments resulting from ESAT Region 9 review of the following analytical data:

Site:

Omega Chem OU2

Site Account No.:

09 BC LA02 CAD042245001

CERCLIS ID No.: Case No.:

None

SDG Nos.:

IQB2427, IQC0445, IQC0602, IQC0945, IQC1079, and

IOC1301

Laboratory:

TestAmerica Analytical Testing Corp.

Analysis:

Hexavalent Chromium

Samples:

15 Hydropunch Water Samples (see Case Summary)

Collection Dates:

February 22, March 5, 6, 8, 9, and 12, 2007

Reviewer:

Stan Kott, ESAT/Laboratory Data Consultants

This report has been reviewed by the EPA TOM for the ESAT contract, whose signature appears above.

If there are any questions, please contact Rose Fong (QA Program/EPA) at (415) 972-3812.

Attachment

SAMPLING ISSUES: [X] Yes [] No

Data Validation Report

Case No.: None

SDG Nos.: IQB2427 – IQC1301 Site: Omega Chem OU2

Laboratory: TestAmerica Analytical Testing Corp.

Reviewer: Stan Kott, ESAT/LDC Date: August 23, 2007

I. CASE SUMMARY

Sample Information

SDG IQB2427 Samples: OC2-HPW8A-W-0-365

SDG IQC0445 Samples: OC2-HPW3A-W-0-379 and OC2-HPW3B-W-0-380

SDG IOC0602 Samples: OC2-HPW4A-W-0-382 and OC2-HPW4B-W-0-383

SDG IQC0945 Samples: OC2-HPW6B-W-0-387 and OC2-HPW5A-W-0-388

SDG IQC1079 Samples: OC2-HPW5B-W-0-389, OC2-HPW5B-W-1-390,

OC2-HPW7B-W-0-391, and OC2-HPW8B-W-0-392

SDG IQC1301 Samples: OC2-HPW1A-W-0-394, OC2-HPW1B-W-1-395,

OC2-HPW2A-W-0-396, and OC2-HPW2B-W-0-397

Concentration and Matrix: Low Concentration Water

Analysis: Hexavalent Chromium

SOW: EPA Method 218.6

Collection Date: February 22, March 5, 6, 8, 9, and 12, 2007

Sample Receipt Date: February 22, March 5, 6, 8, 9, and 12, 2007

Preparation Date: February 22, March 5, 6, 8, 9, and 12, 2007 Analysis Date: February 22, March 5, 6, 7, 8, 9, and 12, 2007

Field OC

Field Blanks (FB): Not Provided

Equipment Blanks (EB): Not Provided Background Samples (BG): Not Provided

Field Duplicates (D1): OC2-HPW5B-W-0-389 and OC2-HPW5B-W-1-390

Laboratory OC

Method Blanks: 7B22111-BLK1, 7C05127-BLK1, 7C06150-BLK1,

7C08101-BLK1, 7C09128-BLK1, and 7C12096-BLK1

Associated Samples: Samples listed above

Matrix Spike (MS): IQB2429-01MS2, IQC0444-03MS1, IQC0600-01MS1,

OC2-HPW6B-W-0-387MS1, OC2-HPW8B-W-0-392MS1,

and OC2-HPW1A-W-0-394MS1

Matrix Spike Duplicate (MSD): IQB2429-01MSD2, IQC0444-03MSD1, IQC0600-01MSD1,

OC2-HPW6B-W-0-387MSD1, OC2-HPW8B-W-0-392MSD1,

and OC2-HPW1A-W-0-394MSD1

Analysis: Hexavalent Chromium

Analyte Hexavalent Chromium Sample Preparation Date February 22, March 5, 6, 8, 9, and 12, 2007 Analysis Date February 22, March 5, 6, 7, 8, 9, and 12, 2007

Sampling Issues

The Chain of Custody (COC) record forms for all SDGs did not specify a sample to be used for laboratory quality control (QC). As a result, the laboratory selected a sample for QC analysis. The effect on data quality is not known.

Additional Comments

As directed by the TOM, a Tier 2 validation (i.e., review all QC results and calibrations, minus calculation check) was performed. A Table 1A is not requested.

For the calibration curve established on March 12, 2007, the calculated percent difference (%D) for calibration standards 0.00030 mg/L and 0.001 mg/L are +68 %D and -12 %D, respectively, and exceed the 10% limit. The 10% limit was derived from the ± 10 % limit used in Method 218.6 to determine the linear dynamic range upper limit. The high and low %D indicates that the calibration may not be linear at the low end of the curve. Since the analytical method does not require analysis of a reporting limit (RL) standard to confirm linearity of the calibration curve at the 0.00030 mg/L RL, results greater than 0.0003 mg/L may have a high bias. Affected sample is OC2-HPW2B-W-0-397 (SDG IQC1301).

Definitions of data qualifiers are listed in Table 1B.

This report was prepared in accordance with the following documents:

- Region 9 Standard Operating Procedure 906, Guidelines for Data Review of Contract Laboratory Program Analytical Services (CLPAS) Inorganic Data Packages; and
- USEPA Method 218.6, Determination of Dissolved Hexavalent Chromium in Drinking Water, Groundwater, and Industrial Wastewater Effluents by Ion Chromatography, Revision 3.3, May 1994.

II. VALIDATION SUMMARY

The data were evaluated based on the following parameters:

	<u>Parameter</u>	<u>Acceptable</u>	Comment
1.	Data Completeness	Yes	
2.	Sample Preservation and Holding Times	Yes	
3.	Calibration	Yes	
	a. Initial		
	b. Initial and Continuing Calibration Verificat	tion	
4.	Blanks	Yes	
5.	Laboratory Control Sample (LCS)	Yes	
6.	Duplicate Sample Analysis	Yes	
7.	Matrix Spike Sample Analysis	Yes	
8.	Field Duplicate Sample Analysis	Yes	
9.	Sample Quantitation	Yes	
10.	Overall Assessment	Yes	
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N/A = Not Applicable

III. OVERALL ASSESSMENT OF DATA

All of the method requirements specified in the EPA Method 218.6 have been met. Reported results for hexavalent chromium in all of the samples were appropriately and correctly calculated.

TABLE 1B

DATA QUALIFIER DEFINITIONS FOR INORGANIC DATA REVIEW

The definitions of the following qualifiers are prepared in accordance with the document *USEPA* Contract Laboratory Program National Functional Guidelines for Inorganic Data Review, October 2004.

- U The analyte was analyzed for, but was not detected above the level of the reported sample quantitation limit.
- J The result is an estimated quantity. The associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The result is an estimated quantity, but the result may be biased high.
- J- The result is an estimated quantity, but the result may be biased low.
- R The data are unusable. The sample results are rejected due to serious deficiencies in meeting Quality Control (QC) criteria. The analyte may or may not be present in the sample.
- UJ The analyte was analyzed for, but was not detected. The reported quantitation limit is approximate and may be inaccurate or imprecise.